

Food Safety and Inspection Service Washington, D.C. 20250

DEC 28

Dr. Óscar González Gutiérrez-Solana Chief of the Spanish Inspection Service Subdirection General de Sanidad Exterior y Veterinaria Ministerio de Sanidad Paseo del Prado, 18 28014 Madrid Spain

Dear Dr. Gutiérrez-Solana:

Enclosed is a copy of the Final report of the Food Safety and Inspection Service's (FSIS) March 21 through April 5, 2001, audit of Spain's meat inspection system. We received your October 10, 2001 letter providing comments on the Draft Final report of the same audit, and made the appropriate editorial changes. Your letter has been incorporated into the Final report as Attachment "G."

We appreciate your thorough review of the FSIS audit findings and the immediate corrective actions taken to ensure that meat products exported to the United States meet U.S. import requirements. We also appreciate the steps taken by the Government of Spain to improve the level of government oversight regarding HACCP implementation. If you have any questions regarding the audit or need additional information, please contact me at telephone number (202-720-3781), facsimile number (202-690-4040), or email address (sally.stratmoen@usda.gov).

Sincerely,

Sally Stratmoen, Chief

Equivalence Section

International Policy Staff

Office of Policy, Program Development

and Evaluation

Enclosure

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Luis Esteruelas, Agricultural Counselor, Embassy of Spain, Washington, DC Lloyd Fleck, Minister-Counselor, American Embassy, Madrid Gerry Keily, Counselor (Agriculture) EU Mission to the US Mary Revelt, Minister/Counselor, US Mission to the EU Brussels John Wilson, FAS Area Officer Catherine Otte, FAS John Prucha, ADA, OPPDE Sally Stratmoen, Chief, ES, IPS, OPPDE Ghias Mughal, Chief, International Review Staff, OFO Amy Winton, State Department Country File-Spain (FY01 Audit)

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<u>Clearance:</u>	<u>Initial</u>	<u>Date</u>
Sally Stratmoen, Chief, ES, FSIS, IPS		
Clark Danford, Acting Chief		

AUDIT REPORT FOR SPAIN

March 21 THROUGH April 5, 2001

INTRODUCTION

Background

This report reflects information that was obtained during an audit of Spain's meat inspection system from March 21 through April 5, 2001. The four establishments certified to export meat to the United States were reviewed. All four establishments were conducting processing operations.

The last audit of the Spain's meat inspection system was conducted in June 2000. The same four establishments were reviewed.

The following were major concerns from the previous audit:

- 1. The HACCP plan did not adequately specify critical limit, monitoring procedures and monitoring frequencies performed for each CCP in Establishments 13, 14, 16, and 20. *Corrected.*
- 2. The HACCP plan did not state adequately the procedures that the establishment will use to verify that the plan is being effectively implemented and the frequencies with which these procedures will be performed in Establishments 13, 14, 16, and 20. *Improvements were made but still inadequate*.
- 3. Corrective actions to be followed in response to a deviation from a critical limit were not addressed adequately in the written HACCP plan in Establishments 13, 14, and 20. *Corrected.*
- 4. Both establishment and inspection personnel had been unaware of the requirement for a final review of all documentation pertaining to the monitoring of critical limits for the product included in each shipment eligible for export to the U.S. The auditor explained the requirements for this pre-shipment review in detail; MSC ordered immediate implementation. Corrected except Establishment 16
- 5. MSC inspection officials were not adequately verifying the establishments' HACCP plan for monitoring critical control points, corrective actions, recordkeeping system and verification procedures. The auditor explained in detail; MSC indicated to implement such a system promptly. *Improvement was made but still inadequate*.

FSIS audited Spain's laboratory using equivalent EU Directives (EN 45001 guidelines), and that the previous issues comply with these equivalent guidelines (a, b, and c).

Spain exports only cured pork to the United States. Restrictions are placed on Spanish beef and fresh pork due to presence of foot and mouth disease, rinderpest, hog cholera and scrapie. Spain is considered to have a substantial risk associated with BSE and swine vesicular disease.

During the period of January 1 to December 31, 2000, Spanish establishments exported 589,907 pounds of cured pork to the U.S. Port-of-entry rejections were for Composition/Standard (0.20 %). During the period of January 1 to February 28, 2001, Spanish establishments exported nearly 23,134 pounds of cured pork s to the U.S. There were no rejections at U.S. ports-of-entry.

PROTOCOL

This on-site audit was conducted in four parts. One part involved visits with Spanish national meat inspection officials to discuss oversight programs and practices, including enforcement activities. The second entailed reviewing a selection of inspection records in Spain's meat inspection headquarters preceding the on-site visits. The third was an on-site visit to each exporting establishment. The fourth was an on-site visit to one government laboratory, which performs analytical testing of field samples for the national residue testing program and performs the presence of microbiological contamination with *Listeria*. The fifth was an on-site visit to autonomous government laboratory, which performs analytical testing of field samples for the national residue testing program and performs the presence of microbiological contamination with *Listeria*, *Salmonella* and *E.coli*.

Program effectiveness determinations focused on five areas of risk: (1) sanitation controls, including the implementation and operation of Sanitation Standard Operating Procedures (SSOPs), (2) animal disease controls, (3) residue controls, (4) slaughter/ processing controls, including the implementation and operation of Hazard Analysis and Critical Control Point (HACCP) systems, and (5) enforcement controls, including the testing program for *Salmonella* species and *listeria*. Spain's inspection system was assessed by evaluating these five risk areas.

During all on-site establishment visits, the auditor evaluated the nature, extent, and degree to which findings impacted on food safety and public health, as well as overall program delivery. The auditor also determined if establishment and inspection system controls were in place. Establishments that do not have effective controls in place to prevent, detect and eliminate product contamination/adulteration are considered unacceptable and therefore ineligible to export products to the U.S., and are delisted accordingly by the country's meat inspection officials.

RESULTS AND DISCUSSION

Summary

All four Establishments (13, 14, 16, and 20) certified to export to the United States were audited; one Establishment (Est.14) was judged Acceptable Subject to Re-review on the next audit. Three Establishments (13,16, and 20) were evaluated as acceptable. Details of audit findings, including compliance with HACCP, SSOPs, and testing programs for *Salmonella* and *listeria* are discussed later in this report.

As stated above, five major concerns had been identified during the last audit of the Spanish meat inspection system, conducted in June 2000. Implementation of a Hazard Analysis-Critical Control Point (HACCP) programs were inadequate in Establishments 13, 14, 16, and 20. Monthly supervisory audits were not made as required by FSIS in Establishments 13, 14, 16, and 20.

During this new audit, the auditor determined that the major concerns had been addressed and improvements were made in Establishments 13, 14, 16, and 20. However, deficiencies regarding implementation of the required HACCP programs still exist in the four establishments visited (Ests. 13, 14, 16, and 20). Details are provided in the Slaughter/Processing Controls section later in this report.

Entrance Meeting

On March 21, 2001, an entrance meeting was held in the Madrid office of the Ministerio De Sanidad Y Consumo (MSC), and was attended by Dr. Oscar Gonzalez Gutierrez Solana, Subdirector General de Sanidad Exterior y Veterinaria; Dr. Jesus Martin Ruiz, Jefe de Area de Veterinaria de Salud Publica; Dr. Margarita Garzon Rigau, Jefe de Servicio Veterinaria Oficial; Dr. Sonsoles Sanchez, Jefe de Area, Ministerio de Agricultura; Mr. Mario Carbajo, Interpretor; Mr. Diego Pazos Moran, Senior Agricultural Specialist, American Embassy; and Dr. Faiz R. Choudry, International Audit Staff Officer, FSIS. Topics of discussion included the following:

- Itinerary and lodging arrangements for the auditor were finalized.
- The auditor shared with the MSC officials the updated data collection instruments for HACCP, *Salmonella* testing, and SSOPs.
- Residue Questionnaire for Spain was discussed.

Headquarters Audit

There had been no changes in the organizational structure or upper levels of inspection staffing since the last U.S. audit of Spain's inspection system in June, 2000.

Prior to the on-site audits of establishments, certain central documents were examined in the office of the meat/poultry inspection headquarters, including the results of the 2000 national residue testing program and the 2001 residue testing plan

- Training records for inspectors and laboratory personnel.
- New laws and implementation documents such as regulations, notices, directives and guidelines.
- Pathogen reduction and other food safety initiatives such as SSOPs, HACCP programs.
- Enforcement records including examples of criminal prosecution, consumer complaints, recalls, seizure and control of noncompliant product, and withholding, suspending, withdrawing inspection services from or delisting an establishment that is certified to export product to the United States.

To gain an accurate overview of the effectiveness of inspection controls, FSIS requested that the reviews of the individual establishments be led by the inspection officials who normally conduct the periodic reviews for compliance with U.S. specifications. The FSIS auditor observed and evaluated the process.

Government Oversight

All inspection veterinarians and inspectors in establishments certified by Spain as eligible to export meat products to the United States were full-time either MSC or Autonomous Government Public-Health employees, receiving no remuneration from either industry or establishment personnel.

Establishment Audits

Four establishments were certified to export meat products to the United States at the time this audit was conducted. All four establishments (13, 14, 16, and 20) were reviewed. In all four establishments, adequate MSC inspection system controls and establishment system controls were in place to prevent, detect and control contamination and adulteration of products. These three establishments were found acceptable. One establishment (Est. 14) was rated acceptable subject to re-review on the next audit because of several deficiencies regarding sanitation and condition of facilities.

Laboratory Audits

FSIS audited Spain's laboratory using equivalent EU Directive (EN 45001 guidelines).

During the laboratory audits, emphasis was placed on the application of procedures and standards that were equivalent to U.S. requirements. Information was also collected about

the risk areas of government oversight of accredited, approved, and private laboratories; intra-laboratory quality assurance procedures, including sample handling; and methodology.

The Institute De Salud Carlos 111, Centro Nacional De Alimentacion Laboratory in Ctra. Majadahonda was audited on March 23, 2001.

Effective controls were in place for sample handling and frequency, timely analysis, data reporting, tissue matrices for analysis, equipment operation and printouts, minimum detection levels, recovery frequency, percent recoveries, check sample frequency, and corrective actions. The methods used for the analyses were acceptable

Spain's microbiological testing program for *Salmonella* and *E.coli* was being performed in the autonomous Gobierno de La Rioja laboratory at the Laboratorio Regional. Consejeria de Agricultura, Ganaderia y Desarrollo Rural in Logrono, was visited on March 30, 2001. Dr. Jose Antonio Garcia Morras is the director at this laboratory. This laboratory is not performing *Salmonella* and *E.coli* testing under FSIS' Pathogen Reduction/HACCP rule but *E.coli* testing is being performed for water and other food products and *Salmonella* for different ready-to-eat products. This laboratory meets the following criteria:

- The laboratory was accredited in May 28, 1999 by Entidad Nacional de Acreditacion (ENAC)/approved by the government.
- The laboratory had properly trained personnel, suitable facilities and equipment, a written quality assurance program, and reporting and record-keeping capabilities.
- Results of analyses were being reported to the government or simultaneously to the government and establishment.
- The autonomous government laboratory uses analytical methods for *Salmonella* (Met/BA/Alim y agua/1), and *E.coli* (Met/BA/Alimentos/4) under the Norma EN 45001-89; UNE 66-501-91 y Guia ISO 25-90

Establishment Operations by Establishment Number

The following operations were being conducted in the four establishments:

Cured/dried pork products - three establishments (13, 14, and 20) Cured/dried chorizos – one establishment (16)

SANITATION CONTROLS

Based on the on-site audits of establishments, Spain's inspection system had controls in place for water potability records; chlorination procedures; back—siphonage prevention; hand washing facilities; sanitizers; separation of operations; pest control and monitoring; temperature control; lighting; work space; ventilation; maintenance and cleaning of over-

product ceilings and equipment; dry storage areas; personal dress, habits, and hygiene; equipment sanitizing; and product handling and storage.

Sanitation Standard Operating Procedures (SSOPs)

Each establishment was evaluated to determine if the basic FSIS regulatory requirements for SSOPs were met, according to the criteria employed in the U.S. domestic inspection program. The data collection instrument used accompanies this report (Attachment A).

Basic Establishment Facilities

• The facility for sanitizing knife in the processing room was designed in such a way that it was not possible to sanitize knife completely and effectively in Establishment 20. *Establishment officials ordered correction*.

Condition of Facilities and Equipment

- Overhead beams and ceilings in the pre-boning cooler over exposed edible product were observed with rust and flaking paint in Establishment 14. *Establishment officials ordered correction immediately*.
- A few working table bottoms and frames for edible product that were not in use but ready for use, in the ham curing and de-boning rooms were found with grease, fat, and black discoloration in Establishment 14. *Establishment officials ordered correction immediately*.
- Band saw, that was not in use but ready for use, in the processing room was found with dried fat and old pieces of meat from previous days' production in Establishment 20. Establishment officials ordered correction immediately.

Cross-Contamination

- Dripping condensate from overhead refrigeration units and pipes that were not cleaned/sanitized was falling onto exposed edible product in the ham de-boning room in Establishment 14. *Neither establishment nor meat inspection officials took corrective action.*
- One employee was observed picking up pieces of strings from the floor and, without washing his hands, handling edible product in the pre-de-boning room in Establishment 14. *Neither establishment nor meat inspection officials took corrective action.*
- An other employee was observed picking up steel from the floor and, without washing his hands or sanitizing his steel, handling edible product in the boning room in Establishment 16. *Neither establishment nor meat inspection officials took corrective action.*

Potential for Cross Contamination

• Hams were contacting employees' working platform at the pre de-boning station in Establishment 14. *Establishment officials took corrective action temporarily and proposed modification to prevent recurrence to GOS inspection officials.*

Reconditioning of contaminated Product

• No facility was provided for dropped ham to be reconditioned in a sanitary manner in the processing room in Establishment 14. *Establishment officials ordered correction immediately*.

ANIMAL DISEASE CONTROLS

Spain's inspection system had controls in place to ensure adequate product identification, restricted product control, and procedures for sanitary handling of returned and rework product. Spain does not have any approved slaughter establishments for export to the United States. All hog carcasses or hams used for exported product to the United States are imported from Denmark and Netherlands. Spain's inspection system had adequate controls in place to ensure control over the above areas, with the following deficiencies. The auditor's findings are presented below for the area of animal disease.

- Containers for edible and inedible product were not identified in the processing room and condemned product was not properly identified and denatured in Establishments 13 and 14. *Establishment officials ordered correction immediately*.
- Containers for edible and inedible product were not identified in the boning and processing rooms in Establishment 16. *Establishment officials ordered correction immediately*.
- Condemned product was not denatured in Establishment 20. *Establishment officials ordered correction immediately*.

No outbreaks of animal diseases with public-health significance have been reported since the previous U.S. audit. At the time of audit, Spain had thirty positive cases for Bovine Spongiform Encephalopathy (BSE) and no positive case was found for Foot and Mouth Disease. Spain is considered to have a substantial risk associated with BSE and Swine Vesicular Disease. APHIS <u>has not</u> declared Spain free of Foot and Mouth Disease, Rinderpest, Hog cholera, and Scrapie.

Farm Visit

Mr. Jose Luis Garcia Ferrero' farm (Selecciones Porcinas, S. A, Finca Quinto del Encinar) located in Santa Cruz del Retamar (Toledo) was visited on April 3, 2001. It was a small

swine breeding farm on a sixty hector land with about 1500 sows, boar including market hogs.

A full time private veterinarian makes the diagnosis, prescription and administers the drugs for treatment. Animals are identified by a single earmark, which identifies the farm, as well as a tattooing mark before leaving farm, the month of the birth of the animal and the code for the farm (premises). Medicated feeds are given to sows, boars and young pigs or breeding stock that will not be going to slaughter in the next 75 days. The farm is required to analyze one sample of medicated feed each year to demonstrate the medicated feed is incompliance. Medicated and non-medicated feed combos are separate.

The swine farm that was visited is licensed to store animal drugs on site. Farms must be specifically approved to store animal drugs on the premises. On those farms which are not approved to store drugs, the veterinarian may only prescribe drugs in amounts that can be used immediately. Records are maintained on all animal drugs requiring prescription, which are written in triplicate so that copies can be maintained by the prescribing veterinarian, filed at the farm, provided to the District where the farm is located and provided to the pharmacy/ wholesaler dispensing the drug. The autonomous government veterinarian and sometime even local police cross checks and verify all the prescriptions written or dispensed in the District.

Certificates (affidavits) are issued for every group of animals moving off of the farm, whether to another farm or to slaughter. Any drugs applied to animals within 75 days of slaughter will be recorded on the transportation documents, with a copy of the prescription attached. Drug inventory and use records are maintained and all drugs are controlled in a locked cabinet or refrigerator.

On-site visits by the autonomous veterinarian are scheduled annually to review the record keeping for veterinarian drug use and checks on feedstuffs. The autonomous government schedules on-farm sampling of animals for drug residue at random basis. No samples are scheduled or collected from live animals under this plan in this farm.

The National Program for Residue Control is based on European Community legislation in force related to the ban of hormonal substances (Council Directive 96/22/EC April 1996) and the control of residues on live animals and animal products (Council Directive 96/23/EC of April 1996). These directives have been transposed into Spanish law through the Royal Decree No. 1749 in 1998.

Reporting Positive Results

Though no violations had occurred at the farm visited, the Regional authorities confirmed that violations are followed up on a case-by-case approach depending upon the substance in question. At the farm, the autonomous government will increase inspections but may not take a sample every time. Intensified sampling is statistically based, and if over half of the samples are positive, the entire herd will be destroyed. If the substance is prohibited, there are criminal sanctions resulting in arrest and possible fines/jail.

RESIDUE CONTROLS

Spain's National Residue Testing Plan for 2001 was being followed, and was on schedule. The Spanish inspection system had adequate controls in place to ensure compliance with sampling and reporting procedures and storage and use of chemicals.

SLAUGHTER/PROCESSING CONTROLS

The Spanish inspection system had controls in place to ensure adequate pre-boning trim, ingredients identification, control of restricted ingredients, formulations, packaging materials, processing schedules, processing equipment, and processing records.

Spain does not have any approved slaughter establishment for export to the United States.

HACCP Implementation

All establishments approved to export meat products to the U.S. are required to have developed and implemented a Hazard Analysis Critical Control Point (HACCP) system equivalent to that of the United States. Each establishment's HACCP system was evaluated according to the criteria employed in the U.S. domestic inspection program. The data collection instrument used accompanies this report (Attachment B).

The HACCP programs were audited and found to meet the basic FSIS regulatory requirements with the following exceptions:

- The HACCP plan did not adequately state the procedures that the establishment will use to verify that the plan is being effectively implemented and the frequencies with which these procedures will be performed in Establishments 13, 14, 16, and 20. *Establishments officials indicated that it would be corrected immediately.*
- MSC inspection officials were not adequately verifying the establishments' HACCP plan for monitoring critical control points, and plant verification procedures in Establishments 13, 14, and 16. MSC officials indicated that it would be corrected immediately.
- Corrective actions were not taken in response to a deviation from a critical limit as addressed in the written HACCP plan in Establishments 14. *Establishment officials indicated that it would be corrected immediately.*
- Establishment verification was inadequate, for a final review of all documentation pertaining to the monitoring of critical limits for the product included in each shipment before that shipment leaves the establishment in Establishment 16. *Establishment officials indicated that it would be corrected immediately.*
- The Ministerio de Sanidad y Consumo (MSC) and Autonomus Government Public Health inspection officials did not have formal HACCP training. *The GOS inspection*

officials indicated that they did not have enough manpower to release officials for HACCP training.

Testing for Generic E. coli

All four establishments were not required to meet the basic FSIS regulatory requirements for generic *E. coli* testing because none of the establishment slaughtered meat for export to the United States. All hams intended for export to the United States were imported from Denmark and Netherlands from slaughter establishments approved to export to the United States. Hog carcasses and/or hams received from domestic slaughter establishments were used for Spanish domestic consumption and/or exported to EU countries.

Additionally, establishments had adequate controls in place to prevent meat products intended for Spanish domestic consumption from being commingled with products eligible for export to the United States.

Control of Listeria monocytogenes

In response to the auditor's inquiry regarding the Spanish establishment officials' evaluation of their HACCP programs to address the risk of *Listeria monocytogenes*, the meat inspection officials provided this information. All four establishments either 1) did not conduct a hazard analysis for *Listeria monocytogenes* to determine the food safety hazards reasonably likely to occur in the production process for ready-to-eat products, or 2) did not have scientific evidence to demonstrate that controls were not needed.

Establishment 14 voluntarily initiated four *Listeria monocytogenes* samples every fifteen days for ready-to eat (RTE) product and ten samples every fifteen days for environment contamination. All product from Establishment 13 is shipped to Establishment 14 for further processing and *Listeria monocytogenes* controls. Establishment 20 is planing to initiate *Listeria monocytogenes* sampling voluntarily in the near future. Establishment 16 was taking six samples voluntarily for each *Listeria monocytogenes* and *salmonella* for ready-to-eat products.

MSC meat inspection officials were taking one sample per month from each establishment for *Listeria monocytogenes* testing on raw product only.

ENFORCEMENT CONTROLS

<u>Inspection System Controls</u>

The MSC inspection system controls [control of restricted product and inspection samples, processed meat reinspection, shipment security, including shipment between establishments, prevention of commingling of product intended for export to the United States with domestic product, monitoring and verification of establishment programs and controls, inspection supervision, and documentation, and the importation of only eligible meat products from

other counties (i.e., only from eligible countries and certified establishments within those countries), for further processing] were in place and effective in ensuring that products produced by the establishment were wholesome, unadulterated, and properly labeled. In addition, adequate controls were found to be in place for security items, shipment security, and products entering the establishments from outside sources.

Testing for Salmonella Species

None of the four establishments reviewed was required to meet the basic FSIS regulatory requirements for *Salmonella* testing because none slaughtered meat for export to the United States. Establishments 13, 14, and 20 were producing dry-cured hams and Establishment 16 was producing dry-cured chorizos. *Salmonella* testing was being conducted on ready-to-eat products in Establishments 14 and 16.

Species Verification Testing

At the time of this audit, Spain was not exempt from the species verification-testing requirement. The auditor verified that species verification testing was being conducted in accordance with FSIS requirements.

Monthly Reviews

The internal audits in Spain were being conducted in three parts as follows:

A) Federal Government; two audits per year by Drs. Margarita Garzon and Julio Navarro, both of whom were veterinarians in the Ministerio de Sanidad y Consumo, under the direct supervision of the Subdirector General de Sanidad Exterior y Veterinaria, Dr. Oscar Gonzalez Gutierrez Solana.

No specific method was used for selecting the review dates of the establishments, but the dates varied from year to year. The internal audit program was applied only to export establishments. The internal audits were conducted once a year, and were announced to the inspection personnel about two weeks in advance. Copy of each internal audit report was kept in the headquarters of the Ministerio de Sanidad y Consumo in Madrid.

- B) Autonomus Government Public Health, one audit per year by a staff veterinarian during any time of the year. A copy of the audit report was kept in the Autonomus Government Public Health office and also in the establishment.
- C) Provincial Government, ten audits per year by a staff veterinarian. No specific method was used for selecting the review dates of the establishments, but the dates varied from each audit. One copy of each internal audit report was kept in the Provincial headquarters and also in the establishments. They were being maintained on file for a minimum of 3 years.

The internal review program was applied only to export establishments. The internal audits were conducted mostly once in two months, and were announced to the inspection personnel, about two weeks in advance; the establishment officials were not informed in advance. The records of audited establishments were kept in the inspection offices of the individual establishments, and copies were also kept in the provincial office.

If an establishment failed to comply with U.S. requirements during an internal audit, it would be immediately delisted for U.S. export. Before it may again qualify for eligibility to be reinstated, MSC meat inspection officials are empowered to conduct an in-depth review, and the results are reported to Dr. Oscar Gonzalez Gutierrez Solana, Subdirector General de Sanidad Exterior y Veterinaria, for evaluation. He would formulate a plan for corrective actions and preventive measures.

Enforcement Activities

Dr. Oscar Gonzalez Gutierrez Solana, Subdirector General, MSC, indicated that Royal Decree 1904/1993 is Spain's legislation to enforce noncompliance when an establishment had not met the regulatory requirements. Under this decree, MSC may temporarily withhold the marks of inspection from specific products, suspend inspection, or withdraw a grant of inspection if an establishment is not meeting crucial requirements.

Exit Meetings

An exit meeting was conducted in Madrid on April 4, 2001. The participants were Dr. Oscar Gonzalez Gutierrez-Solana, Subdirector General de Sanidad Exterior y Veterinaria, MSC; Dr. Margarita Garzon Rigau, Jefe de Servicio Veterinaria Oficial; Dr. Jose L. Paramio Lucas, Jefe de Area, Ministerio de Agricultura; Dr. Juan Jose Martinez, Autonomous Government Public Health Logrono; Dr. Visitacion Cortes Ibanez, Autonomous Government Public Health Vallencia; Mrs. Julia Navardro Pedales, Tecnico, MSC; Dr. Antonio Garcia Jane, Jafe de Seccion de Hygiene Alimentaria, Autonomous Government Castilla-La Mancha; Dr. Jose Juan Sanchez Saez, Subdirector General Laboratory (CAN); Dr. Fernando Tovar, Director General (CNL); Mr. Mario Carbajo, Interpretor; Mr. Diego Pazos Moran, Senior Agricultural Specialist, American Embassy; and Dr. Faiz R. Choudry, International Audit Staff Officer, FSIS.

The deficiencies identified were discussed in detail. Dr. Oscar Gonzalez Gutierrez-Solana reinforced the assurances made by the field personnel during and at the conclusions of the on-site reviews of each establishment, and stated that they would ensure prompt compliance with FSIS import requirements.

A second meeting was conducted with European Commission (EC) in Brussels, Belgium on April 5, 2001. The EC participants were Dr. Paolo M. Drostby, EC Expert, Unit E-3, Health and Consumer Protection Directorate-General; Dr. Thomas Eoin Golden, Principal Administrator, Unit D-2 (Biological Risks), Directorate D, Health and Consumer Protection Directorate-General; Ms. Melinda Sallyards, Agricultural Attaché, United States Mission to

the European Union in Brussels; and Dr. Faiz R. Choudry, International audit Staff Officer, FSIS.

During this meeting, the deficiencies listed below were discussed in detail. Dr. Oscar Gonzalez Gutierrez-Solana, Subdirector General de Sanidad Exterior y Veterinaria, MSC, indicated that the Government of Spain would take the necessary steps to ensure that corrective actions and preventive measures, as promised during the audits and exit meetings in the individual establishments, would be implemented.

- The HACCP plan did not state adequately the procedures that the establishment will use to verify that the plan is being effectively implemented and the frequencies with which these procedures will be performed (Establishments 13, 14, 16, and 20).
- MSC inspection officials were not adequately verifying the establishments' HACCP plan for monitoring critical control points, and plant verification procedures (Establishments 13, 14, and 16).
- Corrective actions were not taken in response to a deviation from a critical limit as addressed in the written HACCP plan (Establishments 14).
- Establishment verification was inadequate for a final review of all documentation pertaining to the monitoring of critical limits for the product included in each shipment before it leaves the establishment (Establishment 16).
- Equipment in contact with product such as working tables and band saws, that were not in use but ready for use, in the ham curing and de-boning rooms and processing rooms were found with grease, fat, and black discoloration, dried fat and old pieces of meat (Establishment 14, and 20).
- Cross contamination and insanitary handling of product were observed in Establishments 14 and 16 such as: a) dripping condensate from overhead refrigeration units and pipes that were not cleaned/sanitized daily, was falling onto exposed edible product in the ham boning room; b) one employee was observed picking up pieces of strings from the floor and, without washing his hands, handling edible product in the pre de-boning room; and c) an other employee was observed picking up steel from the floor and, without washing his hands or sanitizing his steel, handling edible product in the boning room.
- Potential for cross contamination of product such as hams was contacting employees' working platform at the pre-de-boning station in Establishment 14.
- Reconditioning of contaminated product such as facility for dropped ham to be reconditioned in a sanitary manner was not provided in the processing room in Establishment 14.
- Basic establishment facility for sanitizing knife in the processing room was designed in such a way that it was not possible to sanitize knife completely and effectively in Establishment 20.

• Containers for edible and inedible product were not identified in Establishments 13, 14, and 16. Condemned product was not properly identified and denatured in Establishment 13, 14, and 20.

CONCLUSION

Four establishments were reviewed: three were acceptable and one was evaluated as acceptable/re-review. The deficiencies encountered during the on-site establishment reviews were adequately addressed to the auditor's satisfaction. The GOS meat inspection officials reinforced the assurances made by the field personnel during and at the conclusions of the on-site reviews of each establishment and stated that they would ensure prompt compliance.

The establishments have made considerable improvements in the implementation of written SSOPs and HACCP programs. However, deficiencies in these areas still exist and, therefore, additional work is needed to fully comply with FSIS requirements.

The Ministerio de Sanidad y Consumo (MSC) and Autonomus Government Public Health inspection officials did not have formal HACCP training. The GOS inspection officials indicated that they did not have enough manpower to release officials for HACCP training.

Dr. Faizur R. Choudry International Audit Staff Officer (signed)Dr. Faizur R. Choudry

ATTACHMENTS

- A. Data collection instrument for SSOPs
- B. Data collection instrument for HACCP programs
- C. Data collection instrument for *E. coli* testing (*not applicable*)
- D. Data collection instrument for Salmonella testing (not applicable)
- E. Laboratory Audit Forms
- F. Individual Foreign Establishment Audit Forms
- G. Written Foreign Country's Response to the Draft Final Audit Report

Data Collection Instrument for SSOPs

Each establishment was evaluated to determine if the basic FSIS regulatory requirements for SSOPs were met, according to the criteria employed in the U.S. domestic inspection program. The data collection instrument contained the following statements:

- 1. The establishment has a written SSOP program.
- 2. The procedure addresses pre-operational sanitation.
- 3. The procedure addresses operational sanitation.
- 4. The pre-operational procedures address (at a minimum) the cleaning of food-contact surfaces of facilities, equipment, and utensils.
- 5. The procedure indicates the frequency of the tasks.
- 6. The procedure identifies the individuals responsible for implementing and maintaining the activities.
- 7. The records of these procedures and any corrective action taken are being maintained on a daily basis.
- 8. The procedure is dated and signed by the person with overall on-site authority.

The results of these evaluations were as follows:

Est.#	1.Written program addressed	2. Pre-op sanitation addressed	tation sanitation surfaces		5. Frequency addressed	6. Responsible indiv. identified	7. Documentation done daily	8. Dated and signed
13			$\sqrt{}$	V	$\sqrt{}$	V	$\sqrt{}$	$\sqrt{}$
14	√	√	V	V	V	V	√	V
16			$\sqrt{}$	V	$\sqrt{}$	V		$\sqrt{}$
20			$\sqrt{}$		$\sqrt{}$	V		\checkmark

Data Collection Instrument for HACCP Programs

Each of the establishments approved to export meat products to the U.S. was required to have developed and implemented a Hazard Analysis – Critical Control Point (HACCP) system. Each of these systems was evaluated according to the criteria employed in the U.S. domestic inspection program. The data collection instrument included the following statements:

- 1. The establishment has a flow chart that describes the process steps and product flow.
- 2. The establishment had conducted a hazard analysis.
- 3. The analysis includes food safety hazards likely to occur.
- 4. The analysis includes the intended use of or the consumers of the finished product(s).
- 5. There is a written HACCP plan for each product where the hazard analysis revealed one or more food safety hazard(s) reasonably likely to occur.
- 6. All hazards identified in the analysis are included in the HACCP plan; the plan lists a CCP for each food safety hazard identified.
- 7. The HACCP plan specifies critical limits, monitoring procedures, and the monitoring frequency performed for each CCP.
- 8. The plan describes corrective actions taken when a critical limit is exceeded.
- 9. The HACCP plan was validated using multiple monitoring results.
- 10. The HACCP plan lists the establishment's procedures to verify that the plan is being effectively implemented and functioning and the frequency for these procedures.
- 11. The HACCP plan's record-keeping system documents the monitoring of CCPs and/or includes records with actual values and observations.
- 12. The HACCP plan is dated and signed by a responsible establishment official.

The results of these evaluations were as follows:

Est.#	1. Flow diagram	2. Hazard analysis	3. All hazards ident- ified	4. Use & users included	5. Plan for each hazard	6. CCPs for all hazards	7. Mon- itorin g is spec- ified	8. Corr. actions are des- cribed	9. Plan valida- ted	10.Ade- quate verific. Proced- ures	11.Ade- quate docu- mentation	12. Dated and signed
13	√	√	√	√	√	V	√1	√	√	√3	√	√
14	√	√	√	√	√	√	√	√2	√	√	√	√
16	√	√	√	√	√	√	√	√	√	√3	√	√
20	V	√	√	V	√	√	√	V	V	√3	√	V

Attachment E NAME OF FOREIGN LABORATORY REVIEW DATE U.S. DEPARTMENT OF AGRICULTURE FOOD SAFETY AND INSPECTION SERVICE INTERNATIONAL PROGRAMS 03/23/2001 Instituto de Salud Carlos 111 Centro Nacional de Alimentacion FOREIGN COUNTRY LABORATORY REVIEW CITY & COUNTRY ADDRESS OF LABORATORY FOREIGN GOV'T AGENCY Ctra. Majadahonda a Pozuelo, km 2 Majadahonda (Madrid), SPAIN Ministerio de Salud y Consumo 28220 Majadahonda, Madrid-Spain NAME OF FOREIGN OFFICIAL NAME OF REVIEWER Mr. Fernando T. Hernandez, Dir.; Dr. Jose J. S. Saaz, Sub-Dir.; Dr. Margarritta Garzon Dr. Faiz R. Choudry, DVM 300 400 500 200 203 800 Residue Code/Name 111 100 923 List. S/V DEVIEW ITCMS

	REVIEW ITEMS	ITEM #													i
	Sample Handling	01		A	٨	A	A	A	٨	A	A	A	A	A	
OURES	Sampling Frequency	02	CODE	A	٨	٨	A	٨	٨	٨	A	A	٨	٨	
SAMPLING PROCEDURES	Timely Analyses	03	JON CO	٨	٨	٨	A	A	A	A	٨	A	A	A	
PLING	Compositing Procedure	04	VALUATION	o	o	o	o	o	o	o	o	o	O	o	
SAM	Interpret Comp Data	05	5	0	o	o	o	o	o	o	o	o	0	o	
	Data Reporting	06		٨	A	A	A	A	A	A	٨	٨	A	A	
	Acceptable Method	07	CODE	٨	٨	A	A	A	A	A	A	A	A	Α_	:
TICAL	Correct Tissue(s)	08		٨	٨	A	A	٨	A	A	A	A	A	٨	
ANALYTICAL PROCEDURES	Equipment Operation	09	ALUATION	٨	٨	A	A	A	٨	A	A	A	o	o	L
	Instrument Printouts	10	EV,	A	٨	A	A	A	A	A	A	A	o	o	
	Minimum Detection Levels	11		A	٨	A	A	A	A	A	A	A	o	o	
S.	Recovery Frequency	12	CODE	A	٨	٨	A	٨	٨	٨	٨	A	0	О	
URAN RES	Percent Recovery	13		٨	٨	٨	٨	٨	٨	٨	٨	A	0	o	
QUALITY ASSURANCE PROCEDURES	Check Sample Frequency	14	ALUATION	٨	A	A	٨	٨	٨	A	A	A	٨	A	
PRO	All analyst w/Check Samples	15	ALU/	A	٨	٨	A	٨	٨	٨	A	٨	٨	٨	
OĞ.	Corrective Actions	16	2	٨	٨	٨	٨	٨	٨	٨	٨	٨	٨	٨	
	International Check Samples	17		٨	٨	٨	٨	٨	٨	٨	٨	A	A	A	
REVIEW PROCEDURES	Corrected Prior Deficiencies	18	EVAL. CODE		٨	A	٨	٨	٨	A	A	٨	٨	٨	
OTHER REVIEW		19	3000 200E												
OT!		20	EVAL												

SIGNATURE OF REVIEWER

DATE



United States
Department of
Agriculture

Food Safety And Inspection Service Technical Service Center

Suite 300, Landmark Center 1299 Farnam Street Omaha, NE 68102

Questions for Auditing Laboratories

<u>General</u>

Name & location of lab:

Laboratorio Regional de la Comunidad Autonoma

de la Rioja. LOGRONO (SPAIN); March 30, 2001

Private or gov't lab?

Autonomous Government Laboratory

How & when was accreditation obtained? May 28, 1999 by Entidad Nacional de Acreditacion (ENAC). (Norma EN 45001-89; UNE 66-501-91 y Guia ISO 25-90)

How & how often is accreditation maintained?

yearly

When and how is payment for analysis provided?

N/A (Govt. Lab)

Are results released before payment is received?

<u>yes</u>

What are the qualifications of the analyst(s) performing the individual tasks within a method? One pharmacist; One chemist; One analyst

What are the qualifications of the direct supervisor of the analyst(s)?

Veterinarian

Methodology for HACCP Salmonella samples (regulatory labs)

Does this lab analyze HACCP Salmonella samples? Yes, but not for HACCP

How are HACCP Salmonella samples received & recorded? Delivered by inspection

Are HACCP Salmonella samples analyzed on the day of receipt? Seven days

What method(s) is used for HACCP Salmonella samples? Met/BA/Alim y agua/1

Is it a qualitative method (i.e. +/- result)?

<u>Yes</u>

Are HACCP ground beef samples analyzed for Salmonella? No, only RTE product

What is the size of the ground beef test portion? <u>RTE product (25 grams)</u>

What buffer (and what volume) is used for:

Peptona salina

Sponge samples for Salmonella?

N/A

Pouttry rinsates for Salmonella?

N/A

Poultry rinsates for generic E. coli? <u>N/A</u>

What analytical controls are used? <u>Positive/negative</u>

Are they employed for each sample set? Yes

How are HACCP E. coli results calculated and/or expressed? Quantitative method

How are E. coli results recorded:

Data sheets/work sheets? work sheets

Log books? Computer

How and to whom are HACCP E. coli results reported? <u>Inspection officials at the H/Q</u>

Are "check" samples periodically used to test the proficiency of the lab and analysts for generic *E. coli* testing? <u>Yes</u>

- 6. For individual analysts or for the lab as a whole? *Individual analysts*
- 7. What species/strains are used? CECT (Spain); & BCR (EU)
- 8. How many samples are analyzed and how often? <u>Two samples in every</u> two month and one sample p/year as a internal audit
- 9. Are both inoculated and uninoculated samples provided to analysts for the proficiency testing? <u>uninoculated samples</u>
- 10. How many colony-forming units (cfu) per gram are inoculated into the proficiency samples provided to analysts? <u>N/A</u>

NOTE: IF YOU HAVE ANY QUESTIONS REGARDING THIS, FEEL FREE TO CALL EITHER VICTOR COOK OR BONNIE ROSE AT 202-501-6022.

U.S. DEPARTMENT OF AGRICULTURE	REVIE	W DATE	ESTABLISHMENT NO. AND N	IAME	CITY			
FOOD SAFETY AND INSPECTION SERVICE INTERNATIONAL PROGRAMS	0373	26/2001	Est. 13		Olias Del Rey			
FOREIGN PLANT REVIEW FORM	03/2	3072001	Navidul, S. A.		COUNTRY SPAIN			
NAME OF REVIEWER	NAME	OF FORE	IGN OFFICIAL	 - ·	EVALUATION			
Faiz R. Choudry, DVM			Garzon & Dr. Antonio Gard	cia Jane	X Acceptable Acceptable/ Re-review Unac	серсавь		
CODES (Give an appropriate code for each r A = Acceptable M = Margina			below) U = Unacceptable	N	= Not Reviewed O = Does not a	obly		
1. CONTAMINATION CONTROL		Cross c	ontamination prevention	28 A	Formulations s			
(a) BASIC ESTABLISHMENT FACILITIES		Equipme	ent Sanitizing	29 A	Packaging materials	56 A		
Water potability records	01 A	Product	handling and storage	30 A	Laboratory confirmation	57 A		
Chlorination procedures	02 A	Product	reconditioning	31 A	Label approvals	58 A		
Back siphonage prevention	03 A	Product	transportation	32 A	Special label claims	59 O		
Hand washing facilities	04 A	(d) €S	TABUSHMENT SANITATION PRO	GRAM	Inspector monitoring	60 A		
Sanitizers	os A	Effectiv	e maintenance program	33 A	Processing schedules	61 A		
Establishments separation	06 A	Preoper	ational sanitation	34 A	Processing equipment	62 A		
Pestno evidence	07 A	Operation	onal sanitation	35 A	Processing records	93 63		
Pest control program	08 A	Waste	disposal	36 A	Empty can inspection			
Pest control monitoring	09 A		2. DISEASE CONTROL		Filling procedures	65 O		
Temperature control	10 A	Animal	identification	37 O	Container closure exam	66 O		
Lighting	11,	Antemo	rtem inspec. procedures	38 O	Interim container handling	67 O		
Operations work space	12 A	Antemo	rtem dispositions	39 O	Post-processing handling			
Inspector work space	13 O	Human	e Slaughter	40 O	Incubation procedures	69 O		
Ventilation	14 A	Postmo	rtem inspec. procedures	1	Process. defect actions - plant			
Facilities approval	15 A	Postmo	rtem dispositions	42 O	Processing control inspection			
Equipment approval	16 A	Conden	nned product control	43 U	5. COMPLIANCE/ECON, FRAUD CONTROL			
(b) CONDITION OF FACILITIES EQUIPMENT	τ	Restrict	ed product control	40	Export product identification	72 A		
Over-product ceilings	17 A	Returne	d and rework product	45 N	Inspector verification	73 A		
Over-product equipment	16		3. RESIDUE CONTROL		Export certificates	76		
Product contact equipment	12	Residue	program compliance	46 O	Single standard	75 A		
Other product areas (inside)	20 A	Sampli	ng procedures	470	Inspection supervision	76 A		
Dry storage areas	21 A	Residu	e reporting procedures	48 O	Control of security items	77 A		
Antemortem facilities	22 O	Approv	al of chemicals, etc.	49 A	Shipment security	78 A		
Welfare facilities	23 A	Storage	e and use of chemicals	50 A	Species verification	79 O		
Outside premises	24 A		I. PROCESSED PRODUCT CONTR	oc	"Equal to" status	60 A		
(c) PROOUCT PROTECTION & HANOLING	 ì	Pre-bo	ning trim	51 A	Imports	81 A		
Personal dress and habits	25 A	Bonele	ss meat reinspection	52 O				
Personal hygiene practices	26 A	Ingredi	ents identification	53 A				
Sanitary dressing procedures	270	Contro	l of restricted ingredients	5 54 A				

REVIEW DATE ESTABLISHMENT NO. AND NAME

FSIS FORM 9520-2 (2/93)

ILS DEPARTMENT OF AGRICUNTING	BEVIE	WDATE	ESTABLISHMENT NO. AND NAM	AF.	Torry		
U.S. DEPARTMENT OF AGRICULTURE POOD SAFETY AND INSPECTION SERVICE INTERNATIONAL PROGRAMS	1			VIC.	CITY Torrijos		
FOREIGN PLANT REVIEW FORM	03/2	27/2001	Est. 14 Navidul S. A.		COUNTRY SPAIN		
NAME OF REVIEWER Faiz R. Choudry, DVM			I IGN OFFICIAL Garzon & Dr. Antonio garcia	Jane	EVALUATION X Acceptable/	cceptab	
CODES (Give an appropriate code for each A = Acceptable M = Margin			below) U = Unacceptable	N :			
1. CONTAMINATION CONTROL	,		ontamination prevention	28	= Not Reviewed 0 = Does not Formulations	apply	
1. CONTAMINATION CONTROL			·	U 29			
(a) BASIC ESTABLISHMENT FACILITIES		Equipm	ent Sanitizing	٨	Packaging materials	56 A	
Water potability records	01 A	Product	handling and storage	30 M	Laboratory confirmation	57 O	
Chlorination procedures	02 A	Product	reconditioning	31 M	Label approvals	58 A	
Back siphonage prevention	03 A	Product	transportation	32 A	Special label claims	59 O	
Hand washing facilities	04 A	(d) E:	STABLISHMENT SANITATION PROGRA	M	Inspector monitoring	60 A	
Sanitizers	05 A	Effectiv	re maintenance program	33 A	Processing schedules	61 A	
Establishments separation	06 A	Preoper	ational sanitation	34 A	Processing equipment	62 A	
Pestno evidence	07 A	Operati	onal sanitation	35 A	Processing records	63 K	
Pest control program	80 A	Waste	disposal	36 A	Empty can inspection		
Pest control monitoring	09 A		2. DISEASE CONTROL	-	Filling procedures		
Temperature control	10 A	Animal	identification	37 O	Container closure exam	65 O 66 O	
Lighting	114	Antemo	ortem inspec. procedures	38 O	Interim container handling		
Operations work space	12 A	Antemo	ortem dispositions	39 O	Post-processing handling		
Inspector work space	13 O	Human	e Slaughter	40 O	Incubation procedures		
Ventilation	14	Postmo	rtem inspec. procedures	410	Process. defect actions plant		
Facilities approval	15 A	Postmo	rtem dispositions	42 O	Processing control inspection	71 _A	
Equipment approval	16 A	Conder	nned product control	43 U	5. COMPLIANCE/ECON, FRAUD CONTR	IOC	
(6) CONDITION OF FACILITIES EQUIPMEN	ıT	Restric	ted product control	40	Export product identification	72 A	
Over-product ceilings	17 U	Returne	ed and rework product	45 A	Inspector verification	73 A	
Over-product equipment	18 M		3. RESIDUE CONTROL		Export certificates	74 A	
Product contact equipment	19 A	Residu	e program compliance	46 O	Single standard	75 A	
Other product areas (inside)	20 M	Sampli	ng procedures	470	Inspection supervision	76 A	
Dry storage areas	21 A	Residu	e reporting procedures	48 O	Control of security items	777 A	
Antemortem facilities	22 O	Approv	al of chemicals, etc.	49 A	Shipment security	78 A	
Welfare facilities	23 A	Storag	e and use of chemicals	SO A	Species verification	79 O	
Outside premises	24 A	1	4. PROCESSED PRODUCT CONTROL		"Equal to" status	80 A	
(c) PRODUCT PROTECTION & HANDUN	G G	Pre-bo	ning trim	S1 A	Imports	B1 A	
Personal dress and habits	25 A	Bonele	ss meat reinspection	52 O		1	
Personal hygiene practices	26 A	Ingredi	ents identification	53 A		1	
Sanitary dressing procedures	270	Contro	l of restricted ingredients	54 A		1	

U.S. DEPARTMENT OF AGRICULTURE FOOD SAFETY AND INSPECTION SERVICE	REVIE	W DATE	ESTABLISHMENT NO. AND NAM	Ε		CITY			
INTERNATIONAL PROGRAMS	03/2	9/2001	Est. 16			Albelda de Iregi	ua		
FOREIGN PLANT REVIEW FORM	05.2	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	Embutidos Palacios S.A			COUNTRY			
NAME OF REVIEWER Faiz R. Choudry, DVM			IGN OFFICIAL Garzon & Dr. Arnaldo Cabello		EVALUATION X Acceptable Acc	ceptable/ Unacc			
CODES (Give an appropriate code for each $M = M$ argin			below) U = Unacceptable	N =	= Not Reviewed	O = Does not a			
1. CONTAMINATION CONTROL		Cross c	ontamination prevention	28 M	Formulations 5				
(a) BASIC ESTABLISHMENT FACILITIES	Equipm	ent Sanitizing	29 A	Packaging materials					
Water potability records	01 A	Product	handling and storage	30 A	Laboratory confin	mation	57 A		
Chlorination procedures	02 O	Product	reconditioning	31 A	Label approvals		58 A		
Back siphonage prevention	03 A	Product	transportation	32 A	Special label clain	ns	59 O		
Hand washing facilities	04 A	(d) ES	TABLISHMENT SANITATION PROGRA	M	Inspector monitor	ring	60 A		
Sanitizers	05 A	Effectiv	e maintenance program	33 A	Processing sched	ules	61 A		
Establishments separation	06 A	Preoper	ational sanitation	34 A	Processing equipo	nent	62 A		
Pestno evidence	07 A	Operati	onal sanitation	35 A	Processing record	ls	63 A		
Pest control program	96 A	Waste	disposal	36 A	Empty can inspection				
Pest control monitoring	09 A		2. DISEASE CONTROL		Filling procedures		65 O		
Temperature control	10 A	Animal	identification	37 O	Container closure	exam	66 O		
Lighting	11,	Antemo	ortem inspec. procedures	38 O	Interim container	handling	67 O		
Operations work space	12 A	Antemo	ortem dispositions	39 O	Post-processing handling				
Inspector work space	13 O	Human	e Slaughter	40	Incubation procedures				
Ventilation	14 A	Postmo	rtem inspec. procedures	410	Process. defect actions plant				
Facilities approval	15 A	Postmo	rtem dispositions	420	Processing contro	ol - inspection	710		
Equipment approval	16 A	Conden	nned product control	43 M	5. COMPLIANCE/EC	CON. FRAUD CONTRO			
(b) CONDITION OF FACILITIES EQUIPMEN	τ	Restric	ted product control	40	Export product id	entification	72 A		
Over-product ceilings	17 _A	Returne	ed and rework product	45 N	Inspector verifica	tion	73 A		
Over-product equipment	16 A		3. RESIDUE CONTROL		Export certificate	s	74 A		
Product contact equipment	19	Residue	e program compliance	46	Single standard		75 A		
Other product areas (inside)	20 _A	Sampli	ng procedures	470	Inspection superv	vision	76 A		
Ory storage areas	21 A	Residu	e reporting procedures	48 O	Control of securit	ty items	77 _A		
Antemortem facilities	22 0	Approv	al of chemicals, etc.	49 A	Shipment securit	У	76 A		
Welfare facilities	23 A	Storag	e and use of chemicals	SO A	Species verificati	on	79 O		
Outside premises	24 A		4. PROCESSED PRODUCT CONTROL		"Equal to" status		80 A		
(c) PRODUCT PROTECTION & HANDLING		Pre-bo	ning trim	S1 A	Imports		81 _A		
Personal dress and habits	25 A	Bonele	ss meat reinspection	52 A					
Personal hygiene practices	26 A	Ingredi	ents identification	53 A					
Sanitary dressing procedures	270	Contro	l of restricted ingredients	54 A					

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U.S. DEPARTMENT OF AGRICULTURE FOOD SAFETY AND INSPECTION SERVICE INTERNATIONAL PROGRAMS	ĺ	W DATE	ESTABLISHMENT NO. AND NAM	ΙE	City				
FOREIGN PLANT REVIEW FORM	04/0	02/2001	Est. 20 Redondo Iglesias S.A.		COUNTRY SPAIN				
NAME OF REVIEWER Faiz R. Choudry, DVM			IGN OFFICIAL Garzon, Dr.Visitacion Cortes		EVALUATION X Acceptable Re-review				
CODES (Give an appropriate code for each $A = Acceptable$ $M = Margin$			below) U = Unacceptable	N =	= Not Reviewed 0 = Does not	acceptab			
1. CONTAMINATION CONTROL		Cross c	ontamination prevention	28 A	Formulations				
(a) BASIC ESTABLISHMENT FACILITIES		Equipm	ent Sanitizing	29 A	Packaging materials				
Water potability records	01 A	Product	handling and storage	30 A	Laboratory confirmation	57 O			
Chlorination procedures	02 A	Product	reconditioning	31 A	Label approvals	58 A			
Back siphonage prevention	03 A	Product	transportation	32 A	Special label claims	59 O			
Hand washing facilities	04 A	(d) Es	STABLISHMENT SANITATION PROGRA	M	Inspector monitoring	60 A			
Sanitizers	05 M	Effectiv	e maintenance program	33 A	Processing schedules	61 A			
Establishments separation	06 A	Preoper	ational sanitation	34 A	Processing equipment	62 A			
Pestno evidence	07 A	Operati	onal sanitation	35 A	Processing records	63 63			
Pest control program	08 A	Waste	disposal	36 A	Empty can inspection				
Pest control monitoring	09 A		2. DISEASE CONTROL		Filling procedures	65 O			
Temperature control	10 A	Animal	identification	37 O	Container closure exam				
Lighting	11,	Antemo	ortem inspec. procedures	38 O	Interim container handling				
Operations work space	12 A	Antemo	ortem dispositions	39 O	Post-processing handling				
Inspector work space	13 O	Human	e Slaughter	40	Incubation procedures				
Ventilation	14 A	Postmo	rtem inspec. procedures	410	Process. defect actions - plant				
Facilities approval	15 A	Postmo	rtem dispositions	42 O	Processing control inspection	1 76			
Equipment approval	16 A	Conden	nned product control	43 U	5. COMPLIANCE/ECON, FRAUD CONT	ROL			
(b) CONDITION OF FACILITIES EQUIPMEN	τ	Restrict	ted product control	40	Export product identification	72 A			
Over-product ceilings	17 A	Returne	ed and rework product	45 N	Inspector verification	73 A			
Over-product equipment	18 M		3. RESIDUE CONTROL		Export certificates	74 A			
Product contact equipment	19 A	Residue	e program compliance	46 A	Single standard	75 A			
Other product areas (inside)	20 A	Sampli	ng procedures	47A	Inspection supervision	76 A			
Dry storage areas	21 A	Residue	e reporting procedures	48 A	Control of security items	"A			
Antemortem facilities	22 O	Approv	al of chemicals, etc.	49 A	Shipment security	76 A			
Welfare facilities	23 A	Storage	e and use of chemicals	SO A	Species verification	79 O			
Outside premises	24 A		4. PROCESSED PRODUCT CONTROL		"Equal to" status	80 A			
(c) PRODUCT PROTECTION & HANDUNG	`	Pre-boo	ning trim	51 A	Imports	61 A			
Fersonal dress and habits	25 A	Bonele	ss meat reinspection	\$2 O					
ersonal hygiene practices	26 A	Ingredi	ents identification	53 A					
Sanitary dressing procedures	270	Contro	l of restricted ingredients	54 A					

SIS FORM 9520-2 (2/93)

[SEAL]
OFFICE OF THE
SECRETARY-GENERAL DIVISION

OF EXTERNAL AND VETERINARY

HEALTH

DIVISION OF THE SECRETARY'S OFFICE GENERAL DIVISION OF PUBLIC HEALTH AND CONSUMPTION

GENERAL DIVISION
OF EXTERNAL AND
VETERINARY HEALTH

10.10.01 004108 SENT

30/JN/AC

As regards your letter of August 9, 2001, received at this Unit on August 17, 2001, in which you sent the report on the auditor's visit for the Spanish inspection of meat and meat products carried out from March 21 to April 5, 2001 by the FSIS inspector, Dr. Faiz R. Choudry, I am attaching comments in response to said report, as well as the measures adopted by the establishments which have been communicated by the health officials of the Autonomous Communities as regards the report.

Madrid, October 10, 2001 THE ASSISTANT DIRECTOR [SIGNATURE] [illegible] González Gutiérrez-Solana

[SEAL]
MINISTRY OF HEALTH AND CONSUMPTION
General Division of Public Health and Consumption

GENERAL DIVISION OF EXTERNAL AND VETERINARY HEALTH

DR. SALLY STRATMOEN, ACTING DIRECTOR INTERNATIONAL POLICY STAFF OFFICE OF POLICY, PROGRAM DEVELOPMENT AND EVALUATION, WASHINGTON D.C.

DIVISION OF THE SECRETARY'S OFFICE GENERAL DIVISION OF PUBLIC HEALTH AND CONSUMPTION

GENERAL DIVISION
OF EXTERNAL AND
VETERINARY HEALTH

COMMENTS REGARDING THE AUDITOR'S REPORT ON THE SPANISH MEAT AND MEAT PRODUCT INSPECTION SYSTEM, FROM THE AUDIT CARRIED OUT BY THE FSIS INSPECTOR BETWEEN MARCH 21 AND APRIL 4, 2001.

SLAUGHTER/PROCESSING CONTROLS

HACCP Implementation:

Point 2. Regarding these comments, on May 16 2001 the supervisors for the meat and meat product inspection system for US exports met in this office to unify supervision criteria and systems for the establishments (a copy of the meeting convocation is enclosed).

In addition, on April 23, 2001, copies were made of the manual "HACCP Regulatory Process for HACCP-Based Inspection Reference Guide" which was provided by the inspector Dr. Choudry during his last inspection visit to Spain, so that those in charge of the Autonomous Communities have as much information possible.

Listeria monocytogenes control

Regarding the last paragraph about the detection of Listeria monocytogenes in fresh meat, a change in the criteria to be analyzed has been carried out, keeping in mind the observations indicated in the FSIS inspector's form during the auditory visit of June 2000, as well as those indicated verbally during the same inspector's visit of March-April 2001 regarding this matter.

For this reason we have informed the Autonomous Communities as well as the Majadahonda National Feed Center that from the month of May on, the criteria for determining the presence of Listeria shall be meat products (a copy of the reports is attached).

DIVISION OF THE SECRETARY'S OFFICE GENERAL DIVISION OF PUBLIC HEALTH AND CONSUMPTION

GENERAL DIVISION
OF EXTERNAL AND
VETERINARY HEALTH

-ENFORCEMENT CONTROLS

Monthly Reviews

A). Normally the people who carry out the visits are Margarita Garzon and Julia Navarro. The number of supervisory visits from the Ministry of Health and Consumption are at least once a year, although we have been doing two per establishment. The reports from these visits are sent through the central services of the Autonomous Communities to the Provincial Delegations and the Official Veterinarian of the Establishment.

Only those visits carried out by the Ministry at the establishments are announced at the Central Services Offices of the Autonomous Communities where the establishments to be visited are located.

Enforcement Activities

2nd line. The applicable legislation is the Royal Decree 1904/1993, of October 29. In additions, those actions may be carried out under the Ministerial Order of April 4 1995, through which technical and sanitary conditions are regulated applicable to the meat and meat product establishments exporting to the USA.

Attachments

D), In the Foreign Plant Review Form for establishment No. 16 under the item "Name of Foreign Official", the name of Arnaldo Cabello is incorrect, because that person works at the Ministry of Agriculture, Fish, and Feed. The correct name is Juan Jose Martinez de Loza.

CONCLUSION

Paragraph three:

In accordance with the indication given in this paragraph, we inform you that two Advanced Technicians from the General Division of Public Health and Consumption (Ministry of Health and Consumption) have

DIVISION OF THE SECRETARY'S OFFICE GENERAL DIVISION OF PUBLIC HEALTH AND CONSUMPTION

GENERAL DIVISION
OF EXTERNAL AND
VETERINARY HEALTH

requested that the seminar "Meat and Poultry Inspection Seminars for International Government Officials" which will be given by the Food Safety and Inspection Service of the United States Department of Agriculture at the University of Puerto Rico from May 19 to the 31 of 2002.

CORRECTIVE MEASURES ADOPTED BY THE HEALTH AUTHORITIES OF THE AUTONOMOUS COMMUNITIES AS A RESULT OF THE FSIS INSPECTOR'S REPORT.

Regarding the deficiencies encountered during the FSIS inspector's visit, the health authorities of the AC's have informed us of the following:

CASTILLA- LA MANCHA

"ESTABLISHMENT 13, CAMPOFRIO ALIMENTACION, S.A. (previously known as NAVIDUL, S.A.) in Olias de Rey (Toledo):

All the items received a grade of acceptable (A) or not inspected (O) except the following points:

-Point 43: Control of rejected products: was graded as M (marginally acceptable), stating, and I quote, "the rejected materials were not identified nor denaturalized. The Official Services of the establishment ordered its immediate correction".

ESTABLISHMENT 14, CAMPOFRIO ALIMENTACION S.A, (previously known as NAVIDUL, S.A.) in Torrijos (Toledo):

All the items received a grade of acceptable (A) or not inspected (O) except the following points:

- -- Item no. 17: Roofing of the area above the products: graded as U (unacceptable).
 - Item no. 18: Equipment on the products: graded as M (marginally acceptable).
 - Item no. 20: Other product areas: graded as M (marginally acceptable)
 - Item no. 28: Prevention of Cross-contamination: graded as U (unacceptable)

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- Item no. 30: Manipulation and storage of products: graded as M (marginally acceptable).
- Item no. 31: Product reconditioning: graded as M (marginally acceptable).
- Item no. 43: Control of rejected products: qualified as U (unacceptable)

The comments made are reproduced here word for word:

- Leakage and condensation from the refrigeration units and the pipes that were not cleaned on a daily basis, falling onto products that were to be consumed, in the de-boning area. Neither the establishment nor the inspection official took any corrective measures.
- In the roof and beams above the head of the refrigerator holding the products prior to being de-boned, rust and pieces of paint were observed hanging over the products which were to be consumed which could have fallen down on top of them. The Official Services of the establishment ordered the immediate correction of this item.
- A few work tables on the ham curing and de-boning area, which had products on them ready to be used, but which had not yet been used, were found to have grease (both fresh and from machinery) of a black color. The Official Services ordered the immediate correction of this item.
- An employee was observed to pick up some cord or string from the floor and, without washing his hands, he proceeded to manipulate the ham in the predeboning area. Neither the establishment nor the Official Services took any corrective action.
- The hams were in contact with the pre-deboning platforms. The Official Services took immediate corrective action, proposing preventive modifications to the Spanish Government Official Inspectors.
- In the curing area there were no facilities for the pieces of ham to be reconditioned in a sanitary manner. The Official Services of the establishment ordered the immediate correction of this item.
- The containers for the consumable and non-consumable products are not identified and the retained materials were not

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appropriately identified and denaturalized. The Official Services of the establishment ordered the immediate correction of this point.

The report issued expresses the following regarding the HACCP programs:

- Establishment no.13:
- Regarding the HACCP plan, the establishment has not adequately established the verification of the Plan, or its implementation and frequency. The Official Services of the establishment indicated that it would be immediately corrected.
 - > Establishment no.14:
- Regarding the HACCP plan, the establishment has not adequately established the verification of the Plan, or its implementation and frequency. The Official Services of the establishment indicated that it would be immediately corrected.
- The corrective actions were not taken in accordance with the deviation from critical limits referenced in writing in the HACCP Plan. The Official Services of the establishment indicated that it would be immediately corrected.

We also reference the fact that at establishment no. 14 the report indicates that:

- There is a potential danger of cross contamination of products with the work platform in the pre-deboning area.
- The reconditioning of cured ham products shall be carried out in a sanitary manner.

Under the item CONCLUSION in the report, the FSIS inspector indicates that three of the four establishments visited have been acceptable except for one (establishment no. 14), which is considered to be acceptable upon revision (marginal).

CORRECTIVE ACTIONS TAKEN

ESTABLISHMENT 13, CAMPOFRIO ALIMENTACION, S.A. (previously known as NAVIDUL, S.A.) in Olias del Rey (Toledo):

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- The establishment has implemented a line item in the HACCP Plan containing the proper verification procedure for the Plan.
- The container with the non-consumable products has been identified. In order to avoid errors that could result in these products being introduced into the food distribution chain, they will be saturated with dye.
- The container with the consumable products has been identified after being purged, inspected, and analyzed.

ESTABLISHMENT 14, CAMPOFRIO ALIMENTACION, S.A. (previously known as NAVIDUL, S.A.) of Torrijos (Toledo):

- The establishment has implemented a line item in the HACCP Plan containing the proper verification procedure for the Plan.
- The container with the non-consumable products has been identified. In order to avoid errors that could result in these products being introduced into the food distribution chain, they will be saturated with dye.
- The container with the consumable products has been identified after being purged, inspected, and analyzed.
- Control of the temperature in the ham storage areas (regarding the item that mentioned that no corrective action had been taken regarding the deviation from the critical limits referenced by the HACCP plan in writing).
 - o The company has modified the HACCP to the critical temperature limit of this storage area.
 - --Condensation in pipes located above the products:
 - -- The company has established a in the cleaning plan an increase in frequency of the cleaning of the pipes.
 - --Likewise, these pipes have been lined with an insulating material to definitively avoid condensation.
 - --Operator who picked up string from the floor and did not wash his hands before continuing to work with product:

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- Implementation by the establishment of sanitary measures for the Plant Chiefs and for the handlers by the Quality Control Department. In addition, two courses on Good Hygiene Practices have been included in the Personnel Training Plan.
- -- Avoid contact by ham with the shoes of the operators on the platform prior to entering the Deboning Area:
 - The establishment has installed a perimeter defense at this platform, which impedes contact between operators' shoes and the hams hung from hooks.

Criteria for action in the event of a ham falling to the floor:

- The establishment has created informational posters.
- The operators have been instructed
- Adequate measures have been established, among others, to specify that a table-container identified as one to be used for hams that have fallen to the floor, which will serve as a storage place until they can be cleaned.

In addition to the actions described above, reforms have been carried out in the establishment with the purpose of avoiding cross contamination regarding the existing problems expressed.

CONSIDERATIONS OF THE OFFICIAL VETERINARY SERVICES OF ESTABLISHMENT NO. 14

- 1. Keeping in mind that the PCC established in the HACCP Plan at establishment no. 14 are temperature, potable water, nitrifiers, oxygen pressure (slicing) and metal detector (slicing), in the documentation written in said establishment, the controls that have been verified "in situ" are registered in a continuous manner, so as to confirm that they have been carried out. Another aspect that is different is that said points are carried out in accordance with the "new work mechanics" mentioned by the aforementioned FSIS inspector.
- 2. Regarding the temperature, corrective measures have been constantly taken, until the point where we have been able to

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install a reliable temperature measuring system. The corrective measure taken was to take "in situ" and by hand the temperature of the chambers and document this on the sheets for the purpose of having the temperature of the chambers under official control. The Official Services have also taken "in situ" temperatures and coordinated these afterwards with the controls monitored by the company. Regarding whether the product with the temperature deviation was exported to the USA, it is very unlikely given the fact that during the months of December, January, February, and March no products have been exported to the USA. (Clarification report of October 17, 2000 attached as Attachment 1).

- 3. We are also sending documentation (as Attachment II) in which we express that, in regards to picking up items off the floor (string, cords, plastic, knives, etc.), which was the subject of a documented complaint (first verbally) to the person in charge of that area, we have put hygiene measures in place. These measures always include sanitation procedures such as washing, disinfecting, and even sterilization.
- 4. The previous point is directly related to the matter discussed in the report, in which it says that an employee picked up some string or cord off the floor and continued working without any corrective measures being taken either by the establishment or by the Official Veterinary Services present at the Inspection, and that according to the FSIS "manual" it is very important for an inspector to act upon any adverse occurrences (for example this one). The Official Veterinary Services declare that they were not present there at that time (even if the deficiency were as important as the manual says), since they were finishing the task of rejecting and intervening in production line 6 and all the product that passed through the last control because of a temporary failure of the metal detector, considering that this was fairly important. Even the least amount of reasoning would tell you that if one were to act in every single instance in which something is detected on the floor (as is documented in this case), in this case if they had been present they would have proceeded in the same way. That is, after the employee had picked up the string he would have been required to go through the sanitation procedures including rinsing, washing with soap, and drying his hands before going back to his work station. Dated documentation is attached regarding what the Veterinarian was doing at that moment, and that it was dealing with the Rejected and Retained labels. (Documentation of this action is attached as Attachment III).

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5. We are attaching some documentation generated by the Official Veterinary Services of the establishment, which reflect the sequence of actions taken regarding ham on the floor (Attachment IV).

CONCLUSION

With this document regarding the "Comments on the FSIS inspector's report", we believe that we have demonstrated the corrective measures taken at establishments 13 and 14 as a consequence of the grades of "unacceptable or marginally acceptable" in the reports dated March 26 and 27 respectively, issued by Dr. Choudy, DVM.

VALENCIA

"As a consequence of the proof carried out by the Official Veterinary Services of the establishment REDONDO IGLESIAS, S.A. located at Utiel (Valencia), I would like to inform you in accordance with the following aspects:

- Adequate function of the utensil sterilizers (knives) in which the water completely covers the blade of same.
- Placement of identifying labels to not use inadequate utensils because of deficiencies in cleaning, maintenance, or others.
- Marking the rejected products with food ink for their identification and elimination from the human consumption distribution chain.
- Withdrawal of by-products with sufficient frequency, with written documentation of their destination.
- Existence of a final microbiological product-analysis program, as well as a fresh product analysis program, which includes testing for Listeria, Salmonella, and E. Coli".

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LA RIOJA

"Regarding the report of the inspector from the Food Safety and Inspection Service of the USA Department of Agriculture on establishment No. 16 EMBUTIDOS PALACIOS, S.A. on March 29, 2001, we would like to share the following comments:

The establishment will be re-evaluating the HACCP plan so that the verification procedures and frequencies will be effective.

Before exporting a meat product to the United States, the establishment will review the production documents for said product to verify that the critical limits have been complied with.

Likewise, the official veterinary services will take charge of the review and verification to ensure that they are completed correctly.

To avoid cross contamination, the following measures have been adopted:

- Identify the training of employees regarding the adoption of proper handling practices.
- The responsible parties at the establishment as well as the veterinary services will observe the steps in the production process and will adopt any relevant corrective measures.

Madrid, October 10, 2001